



"Hernias, hernias everywhere - what to do about them?"

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Conflicts of interest

 Planning Committee & Faculty Disclosure: The Planning Committee and Faculty have no relevant financial relationships with commercial interests to disclose.





Goals/Questions

- What are the types of pediatric hernias, and how common are each?
- Which pediatric hernias should be referred and when?
- What the concerns with each type of hernia?
- How are pediatric hernias repaired?
- What are long-term concerns and expectations?





- History
 - Greek, Hellenistic (323-31 BC)
 terracotta figures¹
 - Treatment radical excision^{2,3}
 - Ramses V (1156-1152 BC),
 mummy with hernia sac⁴
 - 900 BC: Phoenician statue with tight fitting bandages



Fig. I. A. Male terracotta figurine, Paris, Louvre Museum, D 1206. B: Female terracotta figurine, Jena, Friedrich-Schiller-Universitä, Institut für Geschichte der Medizin, Naturwissenschaft und Technik, Ernst-Haeckel-Haus, V 27.





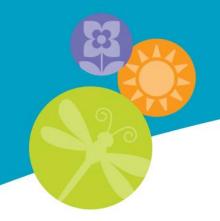


- Galen (200 AD)⁵ & Paul of Aegina (700 AD)⁶ described IH as well
 - Paul recommended ligature of the hernia sac with amputation of the testicle⁷; regression in treatment
- Albucasis (1013-1106 AD) removed testicle during hernia surgery⁸
- William of Salicet (1210-1277 AD) rejected removal of testicle. Described reduction in Trendelenburg⁹





- Ambroise Pare included an entire chapter on hernias in The Apologie and Treatise. Described hernia reduction and closure of peritoneum. Strongly recommended against castration¹⁰
- Kaspar Stromayr distinguished between direct and indirect hernias (1559 AD)
- Hasselbech describes eponymous triangle and iliopubic tract (1814 AD)





- 1870/1871: Lister introduces antiseptic surgery
- 1876: Czerny describes high ligation of the sac
- 1881: Lucas-Championniere splits the external oblique and imbricates the hernia floor
- 1896: Halsted introduces gloves
- Mortality remains high, 2-7%; and recurrence near 100%
- Bassini sutures transversus abdominus and external oblique to inguinal ligament¹¹⁻¹³



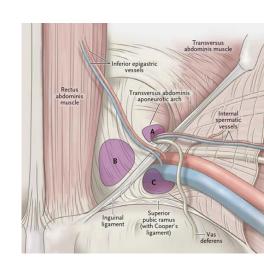


- Background
 - Most common elective pediatric surgical procedure
 - Right 2x > Left
 - More common in premature infants
 - Associated with undescended testicle, abdominal wall defect, connective tissue disease
 - Failure of obliteration of processus vaginalis (testicle, round ligament)





 Direct inguinal hernia bulges through floor medial to inferior epigastric vessels, while indirect hernia bulges lateral to inferior epigastric vessels







- Presentation
 - Asymptomatic or asymptomatic groin swelling that may extend to scrotum or labia
 - Crying or straining makes more prominent
 - Fortunately, most are asymptomatic
 - Rarely, present with incarceration, obstruction or strangulation





- Examination Tips!
 - Stand on contralateral side, palpating along inguinal canal at external ring
 - Valsalva maneuvers (coughing, sit-ups)
 - Although subtle, a thickened cord (silk glove sign) suggests presence of hernia sac
 - Examine for undescended testicles (UDT)
 - Home pictures at end of day









- Diagnosis
 - Physical examination is nearly always all that is needed
 - US if cannot distinguish between hernia and hydrocele, or in rare cases, lymph node
 - Transillumination!





- Incarcerated hernias cannot be reduced. More common in infants.
 - Almost always be reduced with proper technique
 - Straightening canal, <u>constant</u>, <u>directed pressure</u>
 - Trendelenburg position, pain medication and sedation can help
 - If reduced, should be repaired semi-urgently/electively
 - If failed reduction, immediate surgery





- Surgery
 - Do NOT heal spontaneously
 - Must be repaired due to risk of incarceration (17% right, 7% left)
 - Elective repair 11% ED visit by 30 days after dx (ED utilization, HC resources, parental concern)
 - 50% of incarcerations in first 6 months of life, 2/3 in children <1 year





 If asymptomatic, parents do not need to reduce hernia, but should be educated on how to do so, and counseled on signs/symptoms of incarceration





- Timing (AAP Section on Surgery)
 - NICU: 63% repair before discharge, 18% at specific corrected age, 5% when convenient
 - Outpatient: 53% repair when convenient, 27% wait until between 38-60 cGA (mean 53.1 weeks)
- Risk of apnea in premature babies
 - 5% at 48 weeks cGA, 1% at 54 weeks cGA
 - Need 12-24 (preferred) inpatient monitoring earlier





- Operative repair
 - Open
 - High ligation of hernia sac
 - Consider reconstruction of inguinal floor, Bassini repair if weakened



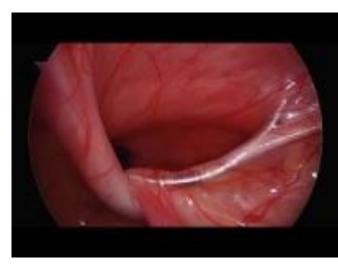








- Operative repair
 - Laparoscopic
 - High ligation of hernia sac without excision
 - Modern studies suggest similar hernia recurrence as with open repair







- Complications
 - Testicular atrophy (1-2%), vas deferens injury (<2%), wound infection (<1%), recurrence (1-5%)
- Follow-up
 - 1 office, phone or televisit
 - 6 months, examine testicles for iatrogenic UDT which would require orchiopexy





- Background
 - Common
 - Gender ratio equal
 - May be more common in African-Americans
 - Increased in Beckwith-Wiedemann syndrome, trisomies 13, 18 and 21







- Presentation
 - Asymptomatic bulge that increases with valsalva
 - Symptoms are rare
 - Obstruction is extremely rare/nearly non-existent
 - No medical treatment (binders/taping are historical interest only)



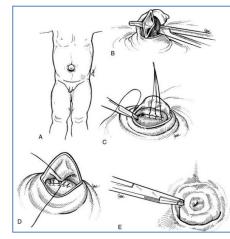


- Surgery
 - Smaller (<1 cm) more likely to close than >1.5 cm
 - No large population studies
 - Most close at 4-5 years of age, 85-90%
 - Theory that transition from abdominal wall to diaphragmatic breathing at this age may contribute
 - Symptomatic repair at time of diagnosis
 - Should repair, will enlarge, risk of complications





- Operative repair
 - Curvilinear incision below umbilicus or vertical through umbilicus
 - Separation of hernia sac from umbilical skin
 - Fascia closed with interrupted absorbable sutures









- Complications
 - Bleeding, infection and recurrence are rare, all ~2%
 - Injury to viscera during repair is very rare
- Follow-up
 - Routine follow-up is offered but not essential
 - Telephone/televisit is reasonable





- Background
 - Hernia through midline/linea alba,
 between xiphoid and umbilicus
 - Small mass, usually with incarcerated preperitoneal fat
 - Different than diastasis recti







- Presentation
 - More often symptomatic, with intermittent pain
 - Midline, epigastric bulge
 - May be adjacent to umbilicus and thus difficult to distinguish from umbilical hernia
 - Congenital, defect in anterior fascia; only preperitoneal fat. Very rarely full thickness, thus almost never incarcerated viscera



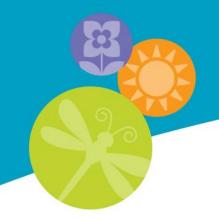


- Surgery
 - Should be repaired because they do not resolve
- Operative repair
 - Mark beforehand! Hard to find when anesthetized
 - If concern for umbilical hernia as well, supraumbilical curvilinear incision permits repair of both
 - Preperitoneal fat excised or reduced, hernia closed





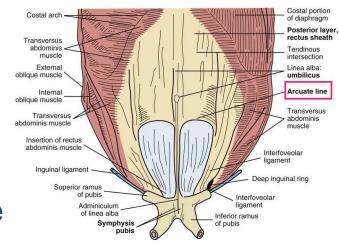
- Complications
 - Bleeding, infection rare
 - Recurrence very rare
- Follow-up
 - Routine follow-up is offered but not essential
 - Telephone/televisit is reasonable





Spigelian Hernia

- Background
 - Defect at junction of rectus
 abdominus and aponeurosis of the
 internal oblique at the linea
 semiluminaris
 - Inferior to arcuate line, where there is no posterior sheath
 - Associated with skeletal anomalies







Spigelian Hernia

- Presentation
 - Pain or a bulge at the lateral border of the rectus muscle, beneath the umbilicus
 - Because the defect is posterior sheath, a bulge may not be obvious, even with incarceration
 - US or CT can be very useful







Spigelian Hernia

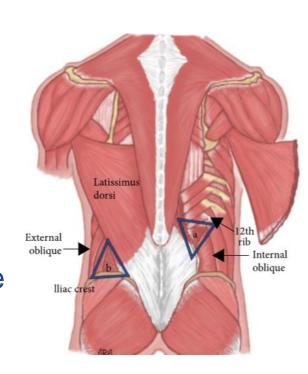
- Surgery
 - Should be repaired because they do not resolve, can become incarcerated
- Operative repair
 - Transverse incision over defect with excision of sac and closure of defect
 - Mesh may be needed in larger defects





Lumbar Hernia

- Background
 - Congenital, very rare
- Presentation
 - Bulge in area bordered by 12th rib,
 sacrospinalis muscle, internal oblique
 - Typically contain preperitoneal fat
 - Soft mass

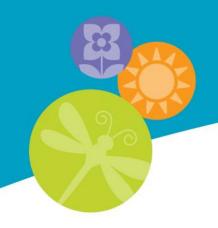






Lumbar Hernia

- Surgery
 - Advised because will not resolve and incarceration is theoretically possible
- Operative repair
 - Primary closure. Prosthetic mesh may be needed because the tissue is often weak or absent when adjacent to bone
 - Absorbable mesh in children, decrease risk of scoliosis

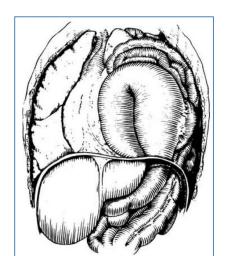


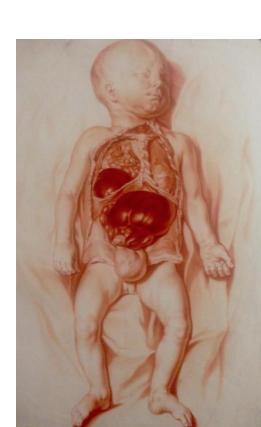


Diaphragmatic Hernia

- History
 - First language description in 1754 by McCauley











Diaphragmatic Hernia

- 1827: Cooper; 1834: Laennec both describe CDH
- Bochdalek (1801-1883) speculates hernia results from rupture of membrane separating the pleuroperitoneal cavity into 2 cavities
- 1888: Naumann; 1890: O'Dwyer make first efforts to repair
- 1901: Aue, first successful repair in adult
- 1905: Heidenhaim, first successful repair in child
- Hedblom's review shows that 75% of 44 infants died





Diaphragmatic Hernia

 1940: Ladd and Gross show consistent, successful repair (9/16 patients)





- Background
 - Incidence around 1/200
 - True incidence likely underestimated due to spontaneous or elective abortion
 - 1/3 die as stillbirth, often with other severe anomalies
 - 80% left sided
 - 90% posterolateral "Bochdalek"; 10% anterior "Morgagni"





- Background
 - Cardiopulmonary disease
 - Pulmonary hypoplasia, with underdeveloped airways, decreased lung weight, decreased number of bronchioles and overall bronchiole cross-sectional area, alveolar count and volume
 - Pulmonary vascularization is abnormal, with resulting pulmonary hypertension





- Diagnosis
 - Prenatal US, most can be seen in 2nd trimester
 - · Polyhydramnios, bowel within chest, MS shift
 - Lung:head ratio (LHR) predicts survival
 - LHR >1.35 100%; 1.35-0.5, 61%; <0.6, no survival
 - LHR should be calculated as observed/expected as the normal ratio changes during gestation
 - Fetal MRI great for morphologic, volumetric measurements





- Presentation
 - Respiratory distress is typical
 - Tachypnea, chest wall retractions, grunting, cyanosis, pallor
 - Scaphoid abdomen
 - CXR confirms bowel within chest







- Management
 - Endotracheal intubation, severe cases (some intubate all)
 - Orogastric decompression
 - Aggressive resuscitation, targeting preductal SaO2 85-95% (cerebral!) with permissive hypercapnea to maintain low PIP
 - ECMO may be needed. CDH accounts for 25% of all infants needing ECMO



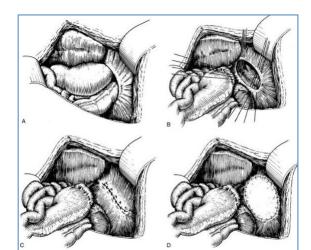


- Surgery
 - Historically performed emergently
 - We now understand that the defect itself is not the issue, but the resulting pulmonary hypoplasia and pHTN.
 Immediate procedures prior to stabilization can exacerbate pHTN and hypoxemia beyond medical correction, resulting in death





- Operative repair
 - Open or minimally invasive from either abdomen or chest
 - Patch may be needed for larger defects









Late-Presenting Diagphragmatic Hernia

- Most often discovered due to non-specific GI or respiratory symptoms prompting work-up
- Complications rare, prognosis favorable
- Symptomatic should be repaired immediately
- Asymptomatic, most would repair electively some would not repair if small and no digestive tract herniation
- Small, right sided defects can avoid repair entirely if covered by liver, especially in high-risk patients





- Inguinal hernia
 - Common. Should be repaired due to the risk of incarceration
 - Outpatient repair should be delayed until 54-60 weeks cGA due to anesthetic risks
 - Open and laparoscopic repairs have similar recurrence
 - Follow-up should focus on testicle location and characteristics in boys





- Umbilical hernia
 - Do not need to be repaired until age 4-5 in the absence of symptoms due to the high rate of spontaneous closure.
 - Incarceration is very rare.
- Epigastric hernia
 - Should be repaired because they do not resolve and are often symptomatic





- Spigelian hernia
 - Very rare
 - Found lateral to rectus muscles and below umbilicus
 - Most present with symptoms at some point, and should be repaired
- Lumbar hernia
 - Very rare
 - Should be repaired due to risk of incarceration





- Diaphragmatic hernia
 - Diagnosed prenatally with US and MRI
 - LHR predicts survival
 - Morbidity and mortality related to pulmonary hypoplasia and pulmonary hypertension
 - Repair in neonatal period when diagnosed
 - Timing of repair of late-presenting CDH should be individualized base on symptoms, risk, and size











Questions?

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- Epic code for internal referrals: St. Vincent REF84BB;
 Clackamas REF84H